



METHOD OF MAKING METALLIC AND/OR CERAMIC PARTS OF NEAR-NET SHAPE

CROSS REFERENCE TO RELATED APPLICATIONS

This application is the US national phase of PCT application PCT/DE2003/003113, filed 19 September 2003, published 13 May 2004 as WO 2004/039748, and claiming the priority of German patent application 10248888.6 itself filed 18 October 2002.

FIELD OF THE INVENTION

The inventions relates to a method of making parts, in particular parts of metallic and/or ceramic powder of near-net shape using the process of powder injection molding (PIM).

STATE OF THE ART BACKGROUND OF THE INVENTION

A) A method of producing metallic parts of near-finish shape.

Metal powder injection molding (MIM) is a method of the mass production of metallic parts, in particular for the production of such parts of near net shape (NNS). The MIM process makes it possible to automatically and inexpensively produce small to medium-large parts in large quantities. The MIM method produces parts with a density equal to 95 to 98% of the theoretical density that can be produced by subsequent hot isostatic compression of the part (without encapsulation).

The method entails the plastification of metal powders with spherical or irregular morphology (particle size from 5 to 300 μm) with a binder system to a so-called feedstock. The goal of the preparation is the coating of all the particles with the organic binder. The feedstock is homogenized in a mixer. Then the

feedstock is charged into the injection-molding machine. In a heated zone components of the binder system (e.g. specific waxes) are melted. A worm advances the thermoplastic mass into the separable mold. After the mold is filled, the liquid mass hardens so the parts can be demolded. The binder system is separated out in a binder-removing step before the sintering. According to the type of binder, additives of different types are also stripped from the part.

There are differences between thermal binder-removing systems (melting out or destruction via the gas phase), solvent extraction, and catalytic solvent-stripping methods. Afterward there is the sintering process in which as a result of diffusion the part is densified up to about 98% of the theoretical density. As a result of the considerable binder content, during sintering there is considerable shrinkage (15 to 20% by volume). Controlling the shrinkage is essential in the production of near-net-shape parts.

Typically used materials for the metallic components in metal-powder injection molding are stainless steel, carbon steel, tool steel, or alloy steel, also ferrite, tungsten carbide, and copper/bronze, cobalt/chromium and tungsten/copper mixtures.

B) Method of making near-net-shape ceramic parts.

The known metal powder injection molding (MIM) method has also been applied to the production of ceramic parts. The so-called powder injection molding (PIM) method also can produce ceramic parts as ceramic injection molding (CIM). To get a corresponding injectable powder mass, organic binder is mixed with

ceramic powder. The injection molding process and the sintering are carried out as in metal-powder injection molding, taking into account the specific characteristics of ceramic powders (e.g. smaller particle size of the starting powder).

5 C) Manufacture of porous metallic parts.

The compression of metal powders to produce porous metal bodies is also known from the literature. To achieve the desired porosity the metal powders can be supplemented with so-called place-holding materials that make it possible to achieve the
10 desired porosity. After compressing the green powder-mixture bodies the place-holding material has to be stripped out of the green bodies so that the green bodies hold nothing but the remaining metal-powder lattice forming empty spaces. The green body thus already has the later porous structure of the shaped
15 body. When the place-holding material is driven out, care is taken to ensure that the metal-powder lattice remains. The subsequent sintering of the green bodies produces a highly porous shaped body, the contacting surfaces of the particles being diffused together when sintered.

20 As place-holding material for forming the porous metallic shaped bodies one uses high-melting-point organic compositions which are vaporized or pyrolyzed (cracked) with release of the thus-produced crack byproducts by means of appropriate solvents from the green bodies. The problem with this is the considerable
25 time it takes to separate the place-holding material and cracking byproducts that react with nearly all the powder-metallurgical metals such as Ti, Al, Fe, Cr, Ni, and the like and that leave

behind high concentrations of impurities. This disadvantage is also encountered when thermoplastics are used, because they have to be heated to get them out of the green body so that the expansion at the glass-transition point is bad for the desired stability of the green body.

In addition the place-holding materials can be inorganic high-melting-point materials such as alkali salts and low-melting metals such as Mg, Sn, Pb, and the like. Such place-holding materials are stripped out in a vacuum or in an inert gas at temperatures of about 600 to 1000°C at considerable cost in terms of energy and time. It is impossible to avoid the harm done by these place-holding materials in particular when used in reactive metal powders such as Ti, Al, Fe, Cr, and Ni.

With alkaline salts there is also the possibility of stripping them out by dissolving them with an appropriate solvent (e.g. water). This method is not ideal for pressed mixtures of metal powders and alkaline salts since the structural integrity of the pressed product is largely lost in the process.

German 196 38 927 describes a method of making highly porous metallic molded bodies wherein first metal powder and a place holder are mixed and then pressed into a green body. Uniaxial and isostatic compression can both be used. The place holder is driven out by heat and the green body is then sintered. If the powder/place-holder mixture is stabilized with a binder, it is usually possible to directly produce a relatively complicated shape in the finished part by multiaxial pressing. The production of a pressing tool to do this is however expensive and difficult.

For small series it is thus advantageous to produce intermediate products with a universal shape (e.g. cylinder or plate) and to transform them by a subsequent mechanical step into the desired end shape.

5 In addition it is necessary to manufacture highly porous parts in large numbers as for use in medicine, air or space travel, or even as filters. Porous parts are made today for example by foaming aluminum or by powder technologies by the use of appropriate place holders. These methods only limitedly allow a
10 near-end shape to be produced of complexly contoured parts in large numbers.

OBJECT AND SOLUTION OF THE INVENTION

It is an object of the invention to produce a method of producing near-net-shape metallic and/or ceramic parts, wherein
15 highly porous parts are produced that have an open porosity of more than 10% by volume, in particular of more than 50% by volume. This object is achieved by a method with all the features of the main claim. Preferred features of the method are seen in the dependent claims.

SUMMARY OF THE INVENTION

20 The present invention relates to the production of complex, highly porous, near-net-shape parts with the so-called powder injection molding (PIM) method. The porosities not to date achievable by this method are attained by the use of certain place
25 holders. With the known powder injection molding (PIM), which includes metal injection molding (MIM) and ceramic injection molding (CIM), metallic and also ceramic powders can be employed.

Above all with metallic powders there is the necessity of leaving minimal residual impurities at the end of the process.

In the following the individual method steps of the production method according to the invention are described with reference to an example.

Starting powder

The above-discussed method is ideal for the use of metallic and also ceramic powders. The particle size of the starting powder can lie in the range of up to 300 μm . To form delicate structures and to achieve a high sintering level between the particles, even finer particles ($< 50 \mu\text{m}$) are preferred. The preferred metallic materials are for example stainless steel, carbon steel, tool steel or alloy steel as well as ferrite, tungsten carbide, and copper/bronze, cobalt/chromium, or even tungsten, copper mixtures.

In addition other starting materials have been tried, in particular titanium, TiAl_6V_4 , nickel-titanium (NiTi) as well as stainless steel (e.g. 316L). These materials have a high affinity for the elements oxygen, carbon, nitrogen, as well as the halogens and thus show what impurities to expect in the sintered parts.

The place holder to use should have the following features:

particle size from 200 to 2000 μm ,

no toxicity,

water-endangerment class 0 to at most 1,

good solubility in water (requirement for stripping out the place holder,

no breakdown up to at most 800°C, and
inexpensive.

It has been discovered according to the invention that in particular NaCl, KCl, K₂CO₃, and Na₂CO₃ fulfill these requirements and give the desired results. These place holders do their jobs particularly well in combination with the MIN feedstock, since the components of the feedstock alone ensure the stability of the parts. It is necessary to take care that the components of the feedstock are not dissolved by the solvent to be used. In order to ensure characteristics reproducible in subsequent parts, the starting characteristics of the employed starting powder (particle distribution, surface composition, chemical analysis of impurities, shake and knock density) must be established.

Production of the injectable mass

The metal powder is plastified and homogenized with the place holder and a binder in a mixer preheated to 110°C. The binder has two components, a plastifier (amide wax) and a component (polyolefin) stabilizing the green starting material.

Shaping injection

The homogenized and reduced mass for injection, comprised of metal powder, place holder, and binder, is plastified in an injection-molding machine at 100°C to 120°C and injected into a mold preheated to between about 30 and 50°C, in particular about 40°C. The pressures and temperatures are set such that there is no separation or nonhomogeneous distribution of the place holder.

Capillary separation of binder from parts (green mass)

The first binder separation leads to an open porosity in the green mass that facilitates in the next method step access of a solvent to the place-holding particles. According to particle size of the starting powder, the binder separation takes place in one or two steps. If the particle size is in the range of 20 to 500 μm a one-stage binder separation is sufficient to achieve the necessary porosity. To this end the green mass is set on a capillary-active material (Al_2O_3 powder) and subjected to a 700 mbar absolute vacuum drying at 130°C. Then the chamber is heated over 2 h to 185°C. This removes about 50% by weight of the fluent component (amide wax). The capillary-active binder separation is also known as "wicking." If the particle size of the starting powder is less than 20 μm , after the capillary first-stage binder separation there is a thermal first-stage binder separation. To do this the parts, after the capillary-active first binder separation, are put in a heated chamber at 20 mbar with an argon throughflow of 10 l/min at 20°C and are heated to 270°C. The heating rate is about 5 K/min. Cooling follows immediately afterward.

Removal of the place holder

After the first-stage binder separation the green mass has a sufficiently open porosity to extract the place holder in the next step at least partially with the aid of a liquid, preferably water. To this end the partially binder-free stable green body is set in a water bath preheated to 40°C to 80°C, the bath being continuously replenished and stirred. When no more ions can be detected, the place holder is almost completely gone. This process

takes for example for a titanium part having a porosity of 70% by volume and an overall volume of 10 cm³ about 3.5 hours.

It is also possible to use any fluid in which the place holder is soluble and that itself does not react chemically with the material of the green body.

Final binder removal of the place-holder-free parts

The removal of the residual binder (generally the stabilizing polyolefin component) is also done thermally. To this end the parts are for example heated in an autoclave at a pressure of 20 mbar with a throughflow of 10 l/min of argon and a heating rate of 4 k/min from room temperature to 480°C, and then immediately cooled and sintered.

Sintering of the place-holder and binder-free sample

According to the material, the parts are subjected to the appropriate sintering process and sintered at temperatures from 800 to 2400°C in the appropriate atmosphere (vacuum, argon, argon/hydrogen, etc). The heating rate and treatment time depend on the workpiece.

The new method according to the invention of powder injection molding is ideal for producing highly porous near-net-shape workpieces from metallic and ceramic starting powders. Complex shapes can be made in large quantities. The place holder used (NaCl, KCl, K₂CO₃, and Na₂CO₃) is not toxic. It can be removed quickly, virtually without residue, and at very low cost (in a water bath) from the parts. Since the stability of the parts is guaranteed by the binder, the parts have their near net shape when

the place holder is removed. The place holder is inexpensive and can, if necessary, be recovered from its solvent.

With the place holder pore sizes in the range of 20 μm to 2 mm and porosities from 10 to 85% by volume, in particular more than 30% by volume or more than 50% by volume can be achieved. The pore distribution is very homogeneous since the feedstock mixes well due to its viscosity. The length of the entire process is mainly determined by the binder removal and the removal of the place holder. Starting with the capillary-active first-stage binder removal and even with the additional expense of a thermal first-stage binder removal (for metal and ceramic powders < 20 μm) the overall time is not more than 14 to 20 hours. This includes the removal of the place holder, the stripping out of residual binder, and the sintering including the heating, cooling, and waiting phases. In addition this is a cost-effective and inexpensive method when medium to large production runs are planned.

Specific Example

In the following the object of the invention is more closely described with reference to an embodiment but without limiting the scope of the invention.

By way of example the method produces a concrete highly porous part (cylinder with a diameter of 25 mm and height of 30 mm when green) from stainless 316L steel (material number 1.4404) with a porosity of at least 70% by volume and pore sizes in the range of 200 to 400 μm . Because of the size of the mixer being used, a charge of 300 cm^3 is produced. The starting materials are:

stainless steel 316L powder (1.4404) with a density of
7.9 g/cm³ and a particle size < 15 µm;
binder 60% by weight of amide wax, trade name C wax
having a density of 1.0 g/cm³ and 40% by weight
polyolefin, trade name PE 520 with a density of 0.93
g/cm³;

place holder of NaCl with density of 2.14 g/cm³ and a
mesh particle size of 200 to 400 µm.

For addition of 70% by volume of place holder and an
overall volume of 300 cm³ the following are used:

470.4 g metal stainless steel 316L powder;
297.3 g NaCl place holder;
59.1 g C wax;
39.4 PE 520.

In order to make the injectable mass the binder is first
melted at 185°C and is then mixed with the metal powder. After
plastifying the metal powder and the binder, the place holder is
added and mixed in for 2 h. The mass is taken out, reduced, and
fed to the injection molder. To work and shape the injectable mass
the following parameters and machine setting are used. Since the
settings depend from the shape of the part as well as the binder
and place-holder content, no absolute values can be given, only
ranges:

mold temperature: 44°C to 49°C;

cylinder zones

Zone 1 (at the nozzle) 144 to 148°C,

Zone 2 135°C,

Zone 3 (input of the mass) 144 to 148°C.

injection pressure 150 to at most 500 bar;

injection time 3 to 6 sec;

injection flow 10 to 50 cm³/sec.

5 Afterward the parts (green) are set on a capillary-active material (Al₂O₃ agglomerate of size 600 to 800 μm). The samples are now set in a vacuum chamber preheated to 130°C. The samples are subject to a vacuum of 700 mbar absolute and preheated in a half hour to 130°C. Afterward they are heated at about 1 K/min to
10 185°C, held for one hour, and then cooled. The samples can be taken out of the vacuum chamber at a temperature of 120°C. Since the particle size of the 316L powder < 15 μm, the samples must be thermally stripped of binder. To this end the samples are heated in a chamber at 20 mbar with an argon throughput at 5 l/min with a
15 heating rate of 4 K/min to 260°C and are then cooled. Subsequently the NaCl place holder is removed in a continuously (magnetically) stirred water bath at 50°C. Gravimetrically the complete removal of the place holder takes 3 h. The samples, which at this stage only contain metal powder, PE 520 wax, and a very small amount of C
20 wax, are heated in a chamber at 20 mbar with an argon throughput of 5 l/min with a heating rate of 5 K/min to 480°C and are then immediately cooled. Gravimetrically the complete removal of all the place holder and binder can be verified. The samples are sitting on an Al₂O₃ base in which the sintering can take place. The
25 sintering is done at 1000°C for 2 h in a vacuum. After the sintering, there is a porosity of 78% by volume. As a result of the reducing during sintering a loss of oxygen and nitrogen can be

demonstrated. The expected increase in carbon content is within acceptable limits.

The drawing shows a microstructure of a highly porous metallic part of 316L that was sintered at 1000°C for 2 h. The pore characteristics are remarkable as they reproduce the cubic crystalline structure of the NaCl particles used as place holder.